

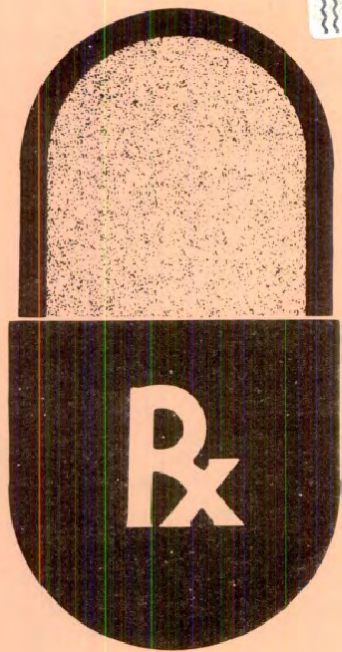
APPROVED DRUG PRODUCTS

With Therapeutic Equivalence Evaluations



The "Orange Book"

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**CUMULATIVE
SUPPLEMENT 3**
AUG'83 - NOV'83

APPROVED PRESCRIPTION DRUG PRODUCTS

WITH
THERAPEUTIC EQUIVALENCE EVALUATIONS

4TH EDITION

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PUBLIC HEALTH SERVICE
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FOOD AND DRUG ADMINISTRATION
APPROVED PRESCRIPTION DRUG PRODUCTS
WITH
THERAPEUTIC EQUIVALENCE EVALUATIONS
CUMULATIVE SUPPLEMENT

I. PREFACE

This cumulative supplement is one of a series of monthly updates to the Approved Prescription Drug Products with Therapeutic Equivalence Evaluations, 4th Edition, (the List) to cover interim revisions to the annual publication of the List in its entirety. The List is comprised of several parts and some by their nature are identified by the term "List". The cumulative supplements routinely provide updates to two of these lists: The Drug Product List and the DESI Addendum.

The List cannot be used effectively without the current cumulative supplement. Users may wish to place an asterisk (*) in the List to the left of the ingredient(s) in the Drug Product List and the product name in the Addendum to indicate that changes to that entry appear in the cumulative supplement. It is also suggested that earlier cumulative supplements be discarded to avoid possible confusion. In this way, only the List and current cumulative supplement need be referenced.

A. DRUG PRODUCT LIST

The Drug Product List cumulative supplements include the changes made since August 1, 1983. Each subsequent cumulative supplement replaces the previous month's cumulative supplement.

Information in this cumulative supplement follows the format of the Drug Product List. The presence of any therapeutic equivalence code indicates that the drug product is multisource; the deletion of a therapeutic equivalence code indicates that the drug product has become single source. (An infrequent exception exists when a therapeutic equivalence code is revised. In that case the deletion of the therapeutic equivalence code is followed immediately by the addition of the revised one.)

Context information on drug products is provided in each cumulative supplement for completeness to assist in locating the proper place in the Drug Product List for the revision. (Strength(s) which already exist in the publication will not be repeated for context.) A page number in parentheses referring to the Drug Product List is located to the right of the ingredient(s).

Additions to the Drug Product List are indicated by new information in the cumulative supplement. Additions new to the current cumulative supplement are indicated by the symbol >_ADD_> to the left of the line on which new information exists. The >_ADD_> symbol is dropped in subsequent cumulative supplements for that item.

Deletions from the Drug Product List are indicated by overstruck print in the cumulative supplement. Deletions new to the current cumulative supplement are indicated by the symbol >DELT (DELETE) to the left of the line containing the overstruck print. The >DELT symbol is dropped in subsequent cumulative supplements for that item.

A newly approved product is identified by the lozenge (◻) to the right of its strength. This identifier remains throughout all cumulative supplements for this edition.

B. ADDENDUM: DESI Pending List

Information in this cumulative supplement follows the format of the Addendum. Additions and deletions are indicated in the same manner as in the cumulative supplement to the Drug Product List. A change in Current Status of a DESI product is also indicated by an addition and a deletion.

II. SPECIAL NOTES

A. REPORT OF COUNTS FOR THE DRUG PRODUCT LIST

Categories of counts derived from product information in the Drug Product List and from this cumulative supplement are presented. The report includes counts of new molecular entities approved by the agency during the current month.

B. PRODUCTS CONTAINING PHENACETIN

The October 5, 1983, Federal Register (48FR45466) provides the following Summary: "The Food and Drug Administration (FDA) is withdrawing approval of new drug applications or parts of new drug applications that provide for drug products containing phenacetin, except for those drug products that are the subject of a hearing request. The basis of the withdrawal is phenacetin's high potential for misuse and its unfavorable benefit-to-risk ratio when incorporated in analgesic combinations which are then subject to excessive chronic use." The effective date of this withdrawal order is November 4, 1983.

Because the subject products are no longer approved, the cumulative supplement will identify them by deleting the applicable active ingredient headers followed by a reference to this Special Note.

III. REPORT OF COUNTS FOR THE DRUG PRODUCT LIST

DESCRIPTION OF REPORT

The following report provides summary counts derived from product information in the Drug Product List and the current cumulative supplement. The counts appear in two sections. Section A. refers to the products in the List and Section B. to products in the current cumulative supplement. A new column of data will appear in Section A. each three-month period following July '83. Section A. therefore will provide baseline and quarterly data while Section B. provides monthly activity.

USE OF REPORT

From the data presented under Section B., users should be able to observe such things as (1) newly approved, DESI effective and remarketed drug products which are added to the List; (2) products that are being removed from the List as the result of withdrawal of approval, changes from prescription to over-the-counter status and discontinued marketing of products; and, (3) trends in approval of products as either multisource or single source during each month within the quarter. The report does not reflect category changes from multisource to single source and vice versa. However, the net gain that results from all additions, deletions and category changes is reflected in the quarterly counts for multisource and single source products.

Drug Product Definition

For this report, a drug product is the representation in the Drug Product List of an active moiety (includes molecular entity and its salts, esters and derivatives) either as a single entity or as a combination product, provided in a specific dosage form and strength for a given route of administration marketed by a firm under a particular generic or trade name.

New Molecular Entity

The active moiety has not previously been approved (either as the parent compound or as a salt, ester or derivative of the parent compound) in the United States for use in a drug product either as a single ingredient or part of a combination.

Drug Product Count

This report provides counts in several categories from the list composed of domestically marketed drug products approved for both safety and effectiveness under sections 505 and 507 of the Federal Food, Drug, and Cosmetic Act. Counts of products still pending in the DESI review are not provided. Excluded also are those approved drug products marketed by distributors; those marketed solely abroad; and products now regarded as medical devices, biologics or foods. Also not included in the counts are those duplicate products of a given applicant whose only distinguishing characteristics are items such as package size, inactive ingredient(s), color and alternate manufacturing sites. These various counts are excluded because the Drug Product List itself excludes products from these categories.

(1) THERAPEUTIC EQUIVALENCE EVALUATIONS PROVIDED ONLY FOR MULTISOURCE PRODUCTS (I.E., AVAILABLE FROM MORE THAN ONE APPLICANT)
 (2) AMINO ACID-CONTAINING PRODUCTS OF VARYING COMPOSITION (SEE PAGE 1-5 OF THE LIST)

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A. COUNTS CUMULATIVE BY QUARTERS		B. ACTIVITY FOR SUPPLEMENT NUMBER 3	
CATEGORIES COUNTED	JULY '83 (BASELINE)	OCT '83	CUMULATIVE
DRUG PRODUCTS LISTED	6679	6783	
SINGLE SOURCE	1908 (28.6%)	1915 (28.2%)	
MULTISOURCE (1)	4771 (71.4%)	4868 (71.8%)	
THERAPEUTICALLY EQUIVALENT	3804 (57.0%)	3891 (57.4%)	
NOT THERAPEUTICALLY EQUIVALENT	957 (14.3%)	967 (14.3%)	
EXCEPTIONS (2)	10 (0.1%)	10 (0.1%)	
NEW MOLECULAR ENTITIES APPROVED	-	2	
NUMBER OF APPLICANTS	304	310	
		B. ACTIVITY FOR SUPPLEMENT NUMBER 3	
		NOV '83	CUMULATIVE
DRUG PRODUCTS ADDED:	43	43	43
NEWLY APPROVED	43	43	43
DESIGN EFFECTIVE	0	0	0
REMARKETED	0	0	0
DRUG PRODUCTS REMOVED:	36	36	36
WITHDRAWN APPROVAL	27	27	27
RX TO OTC SWITCH	0	0	0
DISCONTINUED MARKETING	9	9	9
NET GAIN IN DRUG PRODUCTS	7	7	7
SINGLE SOURCE PRODUCTS APPROVED	10	10	10
MULTISOURCE DRUG PRODUCTS APPROVED	33	33	33
NEW MOLECULAR ENTITIES APPROVED:	3	3	3
AS THE ENTITY	3	3	3
AS A SALT, ESTER OR DERIVATIVE	0	0	0
OF THE ENTITY	0	0	0

REPORT OF COUNTS FOR THE DRUG PRODUCT LIST

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ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE (PAGE 3-2)

TABLET; ORAL
 > DLT > /PERCOCET-S/
 > ADD > PERCOCET
 > DLT > /ACETAMINOPHEN; PHENACETIN; PHENYLPROPANOLAMINE HYDROCHLORIDE/
 > DLT > /PHENYLOXANINE CITRATE/ (PAGE 3-2)

(ALL PRODUCTS - SEE SPECIAL NOTE B.)

ALBUMIN, IODINATED, I-131, SERUM (PAGE 3-4)

INJECTABLE; INJECTION
 > DLT > /RADIOIODINATED SERUM ALBUMIN (HUMAN) I-131/
 > DLT > /MALLINCKRODT/ 16.7-250 MB/ML

AMINOPHYLLINE (PAGE 3-8)

/LIQUID; ORAL/
 SOLUTION; ORAL
 AMINOPHYLLINE
 AA ROXANE LABORATORIES 105MG/5ML

TABLET; ORAL
 AMINOPHYLLINE
 BD BARR LABORATORIES 100MG
 BD 200MG
 AB VANGARD LABS/MHM 100MG
 AB 200MG

/AMPHETAMINE SULFATE/ (PAGE 3-13)

/CAPSULE; CONTROLLED RELEASE; ORAL/
 /BENZEDRINE/
 /SK&F LABORATORIES/ 15MG/
 /TABLET; ORAL/
 /BENZEDRINE/
 /SK&F LABORATORIES/ 5MG/
 /10MG/

AMPICILLIN TRIHYDRATE; PROBENECID (PAGE 3-13)

POWDER FOR RECONSTITUTION; ORAL
 POLYCILLIN-PRB
 > ADD > AB BRISTOL LABS/B-M EQ 3.5GM BASE/100GM/LOT
 > ADD > PROBAMFACIN
 > ADD > AB BIOCRAFT LABS EQ 3.5GM BASE/100GM/LOT

ASPIRIN; BUTALBITAL (PAGE 3-15)

TABLET; ORAL
 AXOTAL
 ADRIA LABORATORIES 650MG;50MG

ASPIRIN; BUTALBITAL; CAFFEINE (PAGE 3-15)

TABLET; ORAL
 ASPIRIN AND CAFFEINE W/ BUTALBITAL
 PUREPAC/KALIPHARMA 325MG;50MG;40MG

> DLT > /ASPIRIN; BUTALBITAL; CAFFEINE; PHENACETIN/ (PAGE 3-15)

(ALL PRODUCTS - SEE SPECIAL NOTE B.)

ASPIRIN; CAFFEINE; DIHYDROCODEINE BITARTRATE (PAGE 3-15)

CAPSULE; ORAL
 SYNALGOS-DC
 IVES LABS/AMMO 356.4MG;30MG;16MG

> DLT > /ASPIRIN; CAFFEINE; PHENACETIN; PROPOXYPHENE HYDROCHLORIDE/
 (PAGE 3-15)

(ALL PRODUCTS - SEE SPECIAL NOTE B.)

ASPIRIN; CAFFEINE; PROPOXYPHENE HYDROCHLORIDE (PAGE 3-16)

CAPSULE; ORAL
 DARVON COMPOUND-65
 AA ELI LILLY INDSTRS/PR 389MG;32.4MG;65MG
 PROPOXYPHENE COMPOUND 65
 AA LEMMON 389MG;32.4MG;65MG
 SK-65 COMPOUND
 AA SK&F LABORATORIES 389MG;32.4MG;65MG

[illegible]

> DLT > / CAFFEINE; CERGOTAMINE; PHENACETIN (PAGE 3-25)

(ALL PRODUCTS - SEE SPECIAL NOTE B.)

CAFFEINE; ERGOTAMINE TARTRATE (PAGE 3-25)

SUPPOSITORY; RECTAL
CAFERGOT
BR SANDOZ PHARMS/SANDOZ 100MG;2MG
MIGRAINE
BR ORGANON/AKZONA 100MG;2MG

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE (PAGE 3-26)

SOLUTION; INTRAPERITONEAL
DIALYTE CONCENTRATE W/
AM MCGAM/AM HOSP DEXTROSE 30% IN PLASTIC CONTAINER
510MG/100ML;30GM/100ML;
200MG/100ML;9.2GM/100ML;
9.6GM/100MLM
510MG/100ML;30GM/100ML;
200MG/100ML;9.4GM/100ML;
11GM/100MLM
DIALYTE CONCENTRATE W/
AM MCGAM/AM HOSP DEXTROSE 50% IN PLASTIC CONTAINER
510MG/100ML;50GM/100ML;
200MG/100ML;9.2GM/100ML;
9.6GM/100MLM
510MG/100ML;50GM/100ML;
200MG/100ML;9.4GM/100ML;
11GM/100MLM

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE (PAGE 3-26)

SOLUTION; INTRAPERITONEAL
DIALYTE W/ DEXTROSE 2.5% IN PLASTIC CONTAINER
AM MCGAM/AM HOSP 26MG/100ML;2.5GM/100ML;
15MG/100ML;560MG/100ML;390MG/100MLM
> ADD >
> ADD >
> ADD >

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM CITRATE (PAGE 3-27)

INJECTABLE; INJECTION
ISOLYTE E IN PLASTIC CONTAINER
AM MCGAM/AM HOSP 35MG/100ML;30MG/100ML;
74MG/100ML;640MG/100ML;
500MG/100ML;74MG/100MLM

CEFTIZOXIME SODIUM (PAGE 3-30)

INJECTABLE; INJECTION
CEFIZOX
SK&F LABORATORIES EQ 1GM BASE/VIALM
EQ 2GM BASE/VIALM

CEFUROXIME SODIUM (PAGE 3-30)

INJECTABLE; INJECTION
ZINACEF
GLAXO EQ 750MG BASE/VIALM
EQ 1.5GM BASE/VIALM

CEPHALOTHIN SODIUM (PAGE 3-31)

INJECTABLE; INJECTION
KEFLIN
ELI LILLY EQ 1GM BASE/VIAL
EQ 2GM BASE/VIAL
> ADD > AP
> ADD > AP
> ADD > AP
SEFFIN
GLAXO EQ 1GM BASE/VIALM
EQ 2GM BASE/VIALM
> ADD > AP
> ADD > AP
> ADD >

CHLORDIAZEPOXIDE (PAGE 3-33)

CAPSULE, CONTROLLED RELEASE; ORAL
LIBREASE
HOFFMANN-LA ROCHE 30MG

CHLORDIAZEPOXIDE HYDROCHLORIDE (PAGE 3-33)

CAPSULE; ORAL
CHLORDIAZEPOXIDE HCL
/ AB / BIDD'S PHARMACEUTICAL / 5MG /
/ AB / / 10MG /
/ AB / / 25MG /

CHLOROTHIAZIDE (PAGE 3-35)

TABLET; ORAL
CHLOROTHIAZIDE
AB CHLSEA LABORATORIES 250MG
AB 500MG

DIATRIZOATE MEGLIUMINE; DIATRIZOATE SODIUM (PAGE 3-57)

INJECTABLE; INJECTION
/DIATRIZOATE MEGLIUMINE AND DIATRIZOATE SODIUM/
DIATRIZOATE-60

DIETHYLPROPION HYDROCHLORIDE (PAGE 3-59)

TABLET; ORAL
DIETHYLPROPION MCL
AA CAMALL 25MG

DIMENHYDRINATE (PAGE 3-60)

INJECTABLE; INJECTION
DIMENHYDRINATE
/AA/ /INJ. MEDICATION SYS/ /15MG/ML/

DIPHENHYDRAMINE HYDROCHLORIDE (PAGE 3-61)

CAPSULE; ORAL
SK-DIPHENHYDRAMINE
/AA/ SK&F LABORATORIES /15MG/

ELIXIR; ORAL
BELTA
AA HALSEY DRUG 12.5MG/5ML
/AA/ /SK-DIPHENHYDRAMINE/ 12.5MG/5ML
/AA/ /SK&F LABORATORIES/ 12.5MG/5ML

DISULFIRAM (PAGE 3-63)

TABLET; ORAL
DISULFIRAM
BX CHELSEA LABORATORIES 250MG
BX 500MG

DOXYCYCLINE HYCLATE (PAGE 3-64)

CAPSULE; ORAL
DOXYCYCLINE HYCLATE
AB CHELSEA LABORATORIES EQ 50MG BASE

INJECTABLE; INJECTION
DOXYCYCLINE
AP ELKINS-SINN EQ 100MG BASE/VIAL
AP EQ 200MG BASE/VIAL
AP VIBRAMYCIN
AP PFIZER LABS/PFIZER EQ 200MG BASE/VIAL

PROPERIDOL; FENTANYL CITRATE (PAGE 3-64)

INJECTABLE; INJECTION
INNOVAR
/JANSSEN PHARMA/ /2.5MG/ML; 0.05MG/ML/
JANSSEN PHARMA 2.5MG/ML; EQ 0.05MG BASE/ML

EPINEPHRINE; LIDOCAINE HYDROCHLORIDE (PAGE 3-66)

INJECTABLE; INJECTION
LIDOCATON
AP PHARMATON/SZ 0.01MG/ML; 122M
AP 0.02MG/ML; 22M

ERGOLOID MESYLATES (PAGE 3-66)

CAPSULE; ORAL
/HYPERGINE/
HYDERGINE LC

ERYTHROMYCIN (PAGE 3-67)

OINTMENT; OPHTHALMIC
ERYTHROMYCIN
AT E FOUGERA/BYK-GLDN 5MG/GM
AT PHARMADERM/BYK-GLDN 5MG/GM
AT XILOTICIN
AT DISTA PRODS/LILLY 5MG/GM

> ADD > ETOPOSIDE (PAGE 3-74)

> ADD > INJECTABLE; INJECTION
> ADD > VEPESID
> ADD > BRISTOL LABS/B-M 20MG/ML

FENTANYL CITRATE (PAGE 3-75)

INJECTABLE; INJECTION
SUBLIMAZE
/JANSSEN PHARMA/ /0.05MG/ML/
JANSSEN PHARMA EQ 0.05MG BASE/ML

FLUOXYMESTERONE (PAGE 3-77)

TABLET; ORAL
FLUOXYMESTERONE
BP COLMED LABORATORIES 10MG

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HYDROCORTISONE (PAGE 3-90)

CREAM; TOPICAL
/AT/ ELDECORT /6.5%/
HYDROCORTISONE
AT BAY LABORATORIES 12M
AT 2.52M

LOTION; TOPICAL
/AT/ /DELACORT/ /0.5%/
/AT/ /MERICON INDUSTRIES/ /0.5%/
/AT/ /GLY-CORT/ /0.5%/
/AT/ /HERAN PHARMACEUTICAL 12M /0.5%/
/AT/ /HYDROCORTISONE/ /0.5%/
/AT/ /MERICON INDUSTRIES/ /0.5%/
/AT/ /TOINE PAULSEN/ /0.5%/

OINTMENT; TOPICAL
HYDROCORTISONE
 AT BAY LABORATORIES 12M
 AT 2.5M
 /SOLUTION/DROPS; OPHTHALMIC/
 /DTEF/
 /UPJOHN/ 10:22/

HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE (PAGE 3-92)

SOLUTION/DROPS; OTIC
NEO-OTOSOL-HC
AT CARTER-GLOGAU LABS 12; EQ 5.5MG BASE/ML; 10,000 UNITS/ML

HYDROCORTISONE ACETATE (PAGE 3-93)

CREAM; TOPICAL
HYDROCORTISONE ACETATE
/6t/ /TONE, PAULSEN/ /12/

LOTION: TOPICAL
PRECORT
 /AT/ INGRAM PHARM 0.5%
 /HYDROCORTISONE "ACETATE"/
 /AT/ TONE PAULSEN /0.5%/
 /SUSPENSION/DROPS/ OPHTHALMIC/ OTIC/
 /HYDROCORTONE/
 /MSD/MERCK/ 1.5%/
 /

HYDROCORTISONE VALERATE (PAGE 3-95)

OINTMENT; TOPICAL
 WESTCORT
 WESTWOOD PHARMS 0.2%

HYDROFLUMETHIAZIDE; RESERPINE (PAGE 3-95)

TABLET; ORAL
HYDROFLUMETHIAZIDE AND RESERPINE
BP COLMED LABORATORIES 50MG:0.125MGX

HYDROXYZINE HYDROCHLORIDE (PAGE 3-96)

	TABLET; ORAL		
	<u>HYDROXYZINE HCL</u>		
> <u>ADD</u> >	<u>AB</u>	BARR LABORATORIES	<u>10MG</u>
	<u>AB</u>	DANBURY PHARMACAL	<u>10MG</u>
	<u>AB</u>		<u>25MG</u>
	<u>AB</u>		<u>50MG</u>

HYDROXYZINE PAMOATE (PAGE 3-96)

CAPSULE; ORAL		
<u>HYDROXYZINE PAMOATE</u>		
AB	VANGARD LABS/MWM	EQ 25MG HCLM
AB		EQ 50MG HCLM

IMIPRAMINE HYDROCHLORIDE (PAGE 3-97)

TABLET; ORAL		
/BP/	/ANTIPRESS/	/ZSNG/
	/LEMON/	
/AB/	/IMAVATE/	/ZSNG/
/AB/	/AK.ROBINS/	/SNG/
<u>THIAPRAMINE HCL</u>		
AB	PAR PHARMACEUTICAL	10MGX
AB		25MGX
AB		50MGX

ISOETHARINE HYDROCHLORIDE (PAGE 3-100)

SOLUTION; INHALATION		
<u>ISOETHARINE HCL</u>		
AN	INTL MEDICATION SYS	0.167%
AN	ROXANE LABORATORIES	0.167%
AN	TRAVENOL LABS	0.25%

ISONIAZID (PAGE 3-101)

SYRUP; ORAL

> ADD >	<u>ISONIAZID</u>	
> ADD > AA	CAROLINA MED PRODS	50MG/5ML
	<u>RIMIFON</u>	
> ADD > AA	HOFFMANN-LA ROCHE	50MG/5ML

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PHENTERMINE HYDROCHLORIDE (PAGE 3-139)

CAPSULE; ORAL
ADIPEX-P
 AA LEMMON 37.5MG
 AA **DAPEX-37.5**
 FERNDALE LABS 37.5MG
 AA **PHENTERMINE HCL**
 CAMALL 37.5MG

POTASSIUM CHLORIDE (PAGE 3-143)

INJECTABLE; INJECTION
POTASSIUM CHLORIDE
 /AA/ TRAVENOL LABS /1MEG/ML/
 /AA/ /3MEG/ML/
 /AP/ /4MEG/ML/

PREDNISOLONE (PAGE 3-145)

TABLET; ORAL
PREDNISOLONE
 /BX/ /REXALL DRUGS/ /5MG/

PREDNISOLONE ACETATE (PAGE 3-146)

INJECTABLE; INJECTION
PREDNISOLONE
 /BP/ /FERNDALE LABS/ /25MG/ML/

PREDNISOLONE SODIUM PHOSPHATE (PAGE 3-147)

SOLUTION/DROPS; OPHTHALMIC
HYDELTRASOL
 /AT/ /MAD/PERCK/ /0.5% PHOSPHATE/
 /AT/ **MEITRON**
 SCHERING EQ 0.5% PHOSPHATE

PREDNISON (PAGE 3-147)

TABLET; ORAL
PREDNISON
 BX DURAMED PHARMS 5MG
 BX 10MG
 BX 20MG

PROCAINAMIDE HYDROCHLORIDE (PAGE 3-149)

CAPSULE; ORAL
PROCAN
 /AA/ PARKE-DAVIS/W-L /144MG/
 /AB/ /300MG/

PROPOXYPHENE HYDROCHLORIDE (PAGE 3-153)

CAPSULE; ORAL
PROPOXYPHENE HCL
 AA RICHLYN LABORATORIES 65MG

PROPYLTHIOURACIL (PAGE 3-154)

TABLET; ORAL
PROPYLTHIOURACIL
 /BD/ /HYLAN PHARMS/ /50MG/

PSEUDOEPHEDRINE HYDROCHLORIDE; TRIPROLIDINE HYDROCHLORIDE (PAGE 3-155)

> ADD > /SYRUP; ORAL/
 SYRUP; ORAL
 /AA/ **PSEUDOEPHEDRINE**
 /AA/ /BAY LABORATORIES/ /30MG/5ML; 1.5MG/5ML/
 /AA/ **TRIPROLIDINE**
 /AA/ /NATL PHARM. INC./ /30MG/5ML; 1.5MG/5ML/
 > ADD > **TRIPOSED**
 > ADD > HALSEY DRUG 30MG/5ML; 1.25MG/5ML

TABLET; ORAL
TRIPROLIDINE
 /AA/ /DANBURY PHARMACAL/ /40MG; 2.5MG/
 > ADD > AA **TRIPROLIDINE AND PSEUDOEPHEDRINE**
 > DLT > /AA/ BOLAR PHARMACEUTICAL 60MG; 2.5MG
 /AA/ **WEST-WARD** /20MG; 2.5MG/
 AA **TRIPROLIDINE HCL AND PSEUDOEPHEDRINE HCL**
 CHLSEA LABORATORIES 60MG; 2.5MG

QUINIDINE SULFATE (PAGE 3-157)

TABLET; ORAL
QUINIDINE SULFATE
 AB VITARINE/WEST CHEM 300MG

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THEOPHYLLINE (PAGE 3-176)

SOLUTION; ORAL
THEOLAIR
 AA RIKER LABS/3M 80MG/15ML
THEOPHYLLINE
 AA ROXANE LABORATORIES 80MG/15MLX
 TABLET, CONTROLLED RELEASE; ORAL
 THEOCONTIN
 BC PURDUE FREDERICK 200MG
THEO-DUR
 BC KEY PHARMACEUTICALS 200MG

THIORIDAZINE HYDROCHLORIDE (PAGE 3-178)

CONCENTRATE; ORAL
THIORIDAZINE HCL
 > ADD > AA CORD LABORATORIES 50MG/MLX
 > ADD > AA 100MG/MLX
 > ADD > AA NATL PHARM MFG/BARRE 100MG/MLX
 TABLET; ORAL
MELLARYL
 AB SANDOZ PHARMS/SANDOZ 100MG
THIORIDAZINE HCL
 > ADD > AB BARR LABORATORIES 10MGX
 > ADD > AB 15MGX
 > ADD > AB 25MGX
 > ADD > AB 50MGX
 > ADD > AB 100MGX
 AB BOLAR PHARMACEUTICAL 10MGX
 AB 100MGX
 AB CORD LABORATORIES 10MGX
 AB 15MGX
 AB 25MGX
 AB 50MGX
 > ADD > AB DANBURY PHARMACAL 10MGX
 > ADD > AB 15MGX
 > ADD > AB 25MGX
 > ADD > AB 50MGX
 > ADD > AB MYLAN PHARMS 100MGX
 AB ZENITH LABORATORIES 100MGX

TIMOLOL MALEATE (PAGE 3-179)

TABLET; ORAL
 BLOCADREN
 MS&D/MERCK 5MG

TRIAMCINOLONE ACETONIDE (PAGE 3-180)

CREAM; TOPICAL
TRIAMCINOLONE ACETONIDE
 AT BAY LABORATORIES 0.025%
 AT 0.1%
 AT 0.5%
 OINTMENT; TOPICAL
TRIAMCINOLONE ACETONIDE
 AT BAY LABORATORIES 0.025%
 AT 0.1%
 AT 0.5%

TRISULFAPYRIMIDINES (PAGE 3-185)

SUSPENSION; ORAL
TRISULFAPYRIMIDINES
 /AB/ /EIDER PHARMS/ /500MG/5ML/
 /AB/ /TRISEN/ /500MG/5ML/
 /AB/ /BEECHAM LABS/BEECHAM/ /500MG/5ML/
 /AB/ /TRISURETO/ /500MG/5ML/
 /AB/ /REID-PROVIDENT LABS/ /500MG/5ML/
 TABLET; ORAL
TRISULFAPYRIMIDINES
 /AB/ /EIDER PHARMS/ /500MG/
 /AB/ /TRIPLE SULFA #2/ /500MG/
 /AB/ /ZENITH LABORATORIES/ /500MG/

TROPICAMIDE (PAGE 3-186)

SOLUTION/DROPS; OPHTHALMIC
MYDRIAFAIR
 AT PHARMAFAIR 0.5%
 AT 1%

TYRANATE (PAGE 3-186)

/CAPSULES; ORAL/
 /TYRANATE/
 /AT ROBINSON/ /250MG/
 /350MG/

VITAMIN A PALMITATE (PAGE 3-188)

CAPSULE; ORAL
VITAMIN A PALMITATE
 /AB/ /TOWNE PHARM/ /60,000 UNITS BASE/

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL;
ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE
HYDROCHLORIDE; RIBOFLAVIN PHOSPHATE SODIUM; THIAMINE
HYDROCHLORIDE; VITAMIN A; VITAMIN E (PAGE AD2)

ADDENDUM

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CURRENT STATUS - INEFFECTIVE

/CARBITAL/ /PARKE-DAVIS/N-1/
/CARBOMAL/ PENTOBARBITAL SODIUM/

/CLISTIN BA/ /MCNEIL PHARM/
/CARBINOXAMINE MALEATE/

/FORHISTAL/ /CIBA/CIBA-GEIGY/
/DIMETHINDENE MALEATE/

> ADD > METHANDROSTENOLONE PAR PHARMACEUTICAL
> ADD > METHANDROSTENOLONE

TUSS-ORNADE SK&F LABORATORIES
CARAMIPHEN EDISYLATE; CHLORPHENIRAMINE MALEATE;
ISOPROPAMIDE IODIDE; PHENYLPROPANOLAMINE HYDROCHLORIDE





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